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Film-laminated flat conductors comprising thermoplastic polyurethanes

5 The present invention relates to film-laminated flat conductors comprising thermoplastic polyurethane and stabilized thermoplastic polyurethane, comprising antioxidants and metal deactivators, preferably for use in film-laminated flat conductors.

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Thermoplastic polyurethane (referred to below as TPU) is an elastomer which is used in many applications, for example shoe applications, films, ski boots and hoses. Cable sheaths constitute a major market for TPU. Here, insulated copper wires

15 are twisted together and then sheathed with TPU. A very wide range of materials, for example PVC or EVA, is used as insulating material for the copper wires. For some applications, TPU is also suitable for the insulation. The sheath in turn is then generally produced from TPU.

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Although the wide use of TPU is based on its good chemical stability, it is advantageous additionally to stabilize the product by introducing additives to prevent damage by heat or UV light. Conventional stabilizers for this purpose are, for

25 example, phenolic antioxidants, hindered amine stabilizers and UV absorbers, various stabilizer classes being used in combination for good UV stabilization.

Furthermore, it is often of interest to reduce the flammability of the TPU and, for example, to produce self-extinguishing cables which do not drip flaming particles. Particularly in cable applications, the incorporation of flame-retardant additives is therefore frequently important. In order to reduce the flammability, flameproofing agents are added to the TPU. Examples

30 of these flame-retardant additives are both compounds which contain halogens, such as chlorine and bromine, and halogen-free flameproofing agents, e.g. phosphoric esters and melamine derivatives. The use of halogen-free flameproofing agents is described in EP-A-617079 and EP-A-189130. The addition of the

35 products can, however, also have an adverse effect on the mechanical properties of the resulting TPU molding materials, for example reduced temperature and heat resistance, especially in the presence of various metals, for example copper.

40 In modern automobiles, electrical consumers and sensors are increasingly being used. As a result, the number of electrical cables required also increases. This leads to increasingly bulky

and complicated cable trees. The increase in weight in the automobile through these cable trees and the susceptibility to faults owing to confusion when connecting cables is, however, undesirable. A development which avoids these disadvantages
5 comprises flat conductors or film-laminated flat conductors.

In this new conductor generation, a plurality of parallel copper strips are insulated from one another or from the environment by two films of plastic. This gives a flat flexible conductor strip
10 which is distinguished by good processing, for example in roofs, low weight, economical production and simple connection.

Owing to their use, for example, in automotive construction, the film-laminated flat conductors have to meet a whole range of
15 requirements with respect to their mechanical, chemical and thermal resistance. It is an object of the present invention to provide a polymer which can advantageously be used for the production of film-laminated flat conductors.

Owing to the design, however, the contact surface between the copper material and the TPU material is particularly large in a film-laminated flat conductor. Moreover, the wall thickness is very small so that destabilization of the TPU by the copper has a considerable effect. It is a further object of the present
20 invention to provide a polymer for the production of film-laminated flat conductors which has high resistance, and in particular high aging resistance, to copper, both when heat is supplied and during storage in water. This also applies in particular when flameproofing agents are added.
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We have found that these objects are achieved by film-laminated flat conductors which are composed of thermoplastic polyurethane which preferably contains an additive mixture comprising an antioxidant (component i) and a metal deactivator (component ii)
30 and, if required, a flameproofing agent (component iii).
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The present invention therefore relates to TPU-containing film-laminated flat conductors and stabilized TPU, comprising (i) antioxidants and (ii) metal deactivators, and the use of the
40 novel TPU for the production of film-laminated flat conductors.

The present invention also relates to a metal deactivator concentrate for the preparation of the novel thermoplastic polyurethanes, comprising thermoplastic polyurethane and metal
45 deactivators (ii) in an amount of from more than 2.5 to 40% by

weight, based on the total weight of the thermoplastic polyurethane.

The present invention furthermore relates to a process for the
5 production of novel film-laminated flat conductors by providing two TPU films and then laminating metal strips between the two films

or

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by coextruding TPU films with metal strips.

The present invention furthermore relates to the use of the novel film-laminated flat conductors in means of transport, machines
15 and/or electrical articles, and means of transport, machines and/or electrical articles comprising the novel film-laminated flat conductors.

Suitable antioxidants which constitute the component (i) in the
20 novel stabilized TPU are in general substances which retard or prevent undesired oxidation processes in the plastic to be protected. In general, antioxidants are commercially available. Examples of antioxidants are sterically hindered phenols, aromatic amines, thiosynergists, organophosphorus compounds of
25 trivalent phosphorus and hindered amine light stabilizers. Examples of sterically hindered phenols are to be found in Plastics Additive Handbook, 5th edition, H. Zweifel, ed, Hanser Publishers, Munich, 2001 ([1]), pages 98-107 and pages 116 - 121. Examples of aromatic amines are to be found in [1], pages
30 107-108. Examples of thiosynergists are given in [1], pages 104-105 and pages 112-113. Examples of phosphites are to be found in [1], pages 109-112. Examples of hindered amine light stabilizers are given in [1], pages 123-136. Phenolic antioxidants are preferably used in the novel mixture of
35 antioxidants.

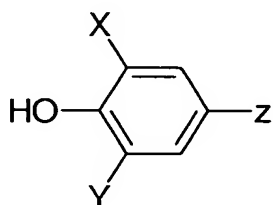
In a preferred embodiment, the antioxidants (i), in particular the phenolic antioxidants, have a molar mass of more than 350, particularly preferably more than 700, g/mol and a maximum molar
40 mass of <10 000, preferably <3 000, g/mol. Furthermore, they preferably have a melting point of less than 180°C, particularly preferably less than 130°C. Moreover, antioxidants which are amorphous or liquid are preferably used. Mixtures of two or more antioxidants may also be used as component (i).

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The abovementioned boundary conditions regarding molar mass and melting point ensure that the antioxidant does not volatilize even in the case of high surface area/volume ratios and that the antioxidant can be uniformly and homogeneously incorporated into the TPU during the synthesis. Examples of suitable phenolic antioxidants are molecules which contain the structure 1 as an active ingredient group.

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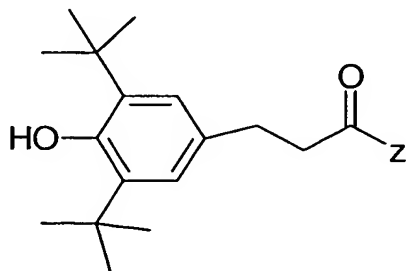
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where X and Y, independently of one another, are a hydrogen atom or straight-chain, branched or cyclic alkyl radicals of 1 to 12 carbon atoms and

20 Z is a covalent bond via which the active ingredient group is bonded to the remaining molecule of the antioxidant (i).

Preferably used phenolic antioxidants (i) are compounds which contain the radical 2.

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35 where Z is as defined above.

Examples of preferred phenolic antioxidants which contain the active ingredient group 1 are triethylene glycol bis(3-(5-tert-butyl-4-hydroxy-m-tolyl)propionate) (Irganox[®] 245, Ciba Spezialitätenchemie AG), hexamethylene bis(3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionate) (Irganox[®] 259), pentaerythrityl tetrakis(3-(3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl)-propionates) (Irganox[®] 1010), octadecyl 3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionate (Irganox[®] 1076), N,N'-hexamethylenebis(3,5-di-tert-butyl-4-hydroxy-hydrocinnamide) (Irganox[®] 1098),

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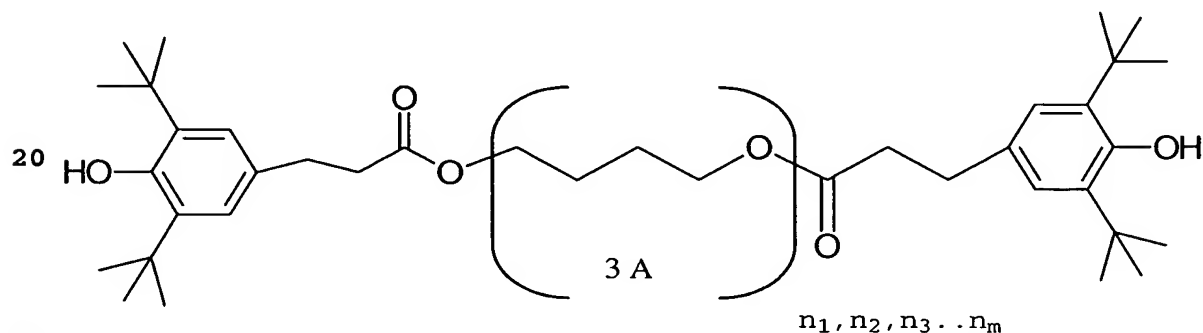
phenol, 2,4-dimethyl-6-(1-methylpentadecyl)+ octadecyl
3-(3,5-di-tert-butyl-4-hydroxy-phenyl)propionate (Irganox®
1141),

2-tert-butyl-6-(3-tert-butyl-2-hydroxy-5-methylbenzyl)-4-methyl-
5 phenyl acrylate (Irganox® 3052) and CAS No. 125643-61-0 (Irganox®
1135).

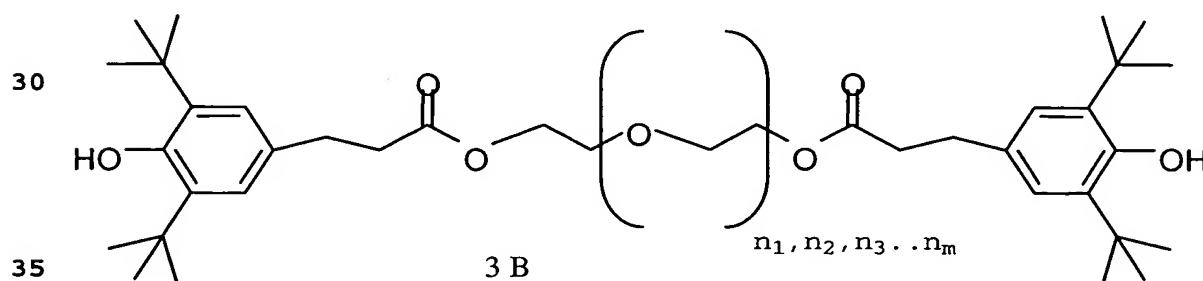
Particularly preferred phenolic antioxidants are Irganox® 1010,
Irganox® 245, Irganox® 259 and antioxidants based on
10 polyetherols.

Antioxidants which can be described by the general formulae 3 A
and 3 B are furthermore particularly preferred.

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In the formulae 3 A and 3 B, $n_1, n_2, n_3 \dots n_m$ are integers > 0 and
the relationships $n_2 = n_1 + 1$, $n_3 = n_2 + 1$, $n_{x+1} = n_x + 1$, etc. apply.

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The preferred antioxidants 3 A and 3 B are accordingly mixtures
of different compounds which differ only in the magnitude of n
and are referred to below as antioxidant mixture. The proportion
of the molecules n_1, n_2, n_3 to n_m is chosen so that the number
45 average molar mass of the antioxidant mixture corresponds to the
molar mass recognized as being advantageous. A proportion of the
molecules n_1, n_2, n_3 to n_m is preferably chosen so that the number

average molar mass of the antioxidant mixture 3 A and/or 3 B is greater than 350, particularly preferably > 700, g/mol and <10 000, preferably <3 000, g/mol.

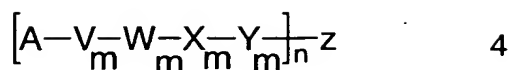
- 5 In a further preferred embodiment, antioxidant mixtures used are those whose polydispersity P_d is greater than 1, i.e. their number average molar mass is less than their weight average molar mass. This is fulfilled, for example, when the antioxidant consists of a mixture of different molecules of the structure 3 A
10 or 3 B with different n.

It may be advantageous if, instead of a single phenolic antioxidant, a mixture of phenolic antioxidants is used for stabilization. In principle, all phenolic antioxidants which
15 comply with the conditions described above with respect to molar mass and melting point can be used for such mixtures.

Mixtures which contain Irganox® 1010 and/or those mixtures which contain phenolic antioxidants according to the formulae 3 A and 3
20 B are particularly preferred.

Metal deactivators are used as component (ii). Metal ions generally catalyze the decomposition of peroxides and thus accelerate the degradation of polymers; consequently, metal
25 deactivators are substances which prevent the harmful effect of metal ions with regard to the degradation of polymers, for example through complexing of these metal ions. Examples of metal deactivators are 2-(2-benzimidazolyl)phenol, 3-(2-imidazolin-2-yl)-2-naphthol, 2-(2-benzoxazolyl)phenol,
30 4-diethylamino-2,2'-dioxy-5-methylazobenzenes, 3-methyl-4-(2-oxy-5-methylphenylazo)-1-phenyl-5-pyrazolones, tris(2-tert-butyl-4-thio(2'-methyl-4'-hydroxy-5'-tert-butyl)phenyl-5-methyl)phenyl phosphite, decamethylenedicarboxydisalicyloylhydrazides,
35 3-salicyloylamino-1,2,4-triazole, 2',3-bis-((3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionyl))propionhydrazide and 2,2'-oxamidobis(ethyl-3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionates). In general, derivatives of hydrazine, such as adipic acid
40 dihydrazide or oxalic acid dihydrazide, and derivatives of oxamide may furthermore act as metal deactivators.

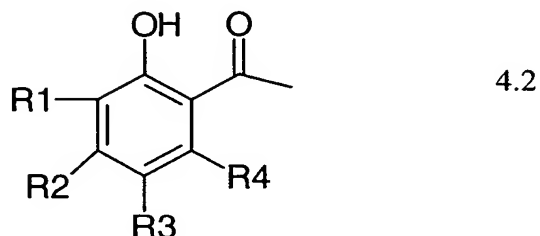
However, it has surprisingly been found that particularly good results are obtained with metal deactivators which are triazole
45 and hydrazine derivatives of salicylaldehyde, for example those metal deactivators which have the structure 4.



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In structure 4, A is a salicylaldehyde radical according to formula 4.2,

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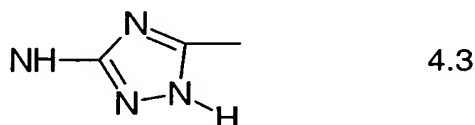


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R1-R4, independently of one another, are H, OH, halogen, NO₂, NH₂, a linear, branched or cyclic C1-C12 alkyl radical, a linear, branched or cyclic C1-C12 alkenyl, a linear, branched or cyclic C1-C12 alkynyl radical or haloalkyl radical, a linear, branched or cyclic C1-C12 alkoxy radical, a linear, branched or cyclic C1-C12 alkylamino radical, containing primary and/or secondary and/or tertiary amines, a linear, branched or cyclic C1-C12 alkylthio radical or a linear, branched or cyclic C1-C12 alkoxy carbonyl radical.

V_m in formula 4 is a triazole radical according to formula 4.3

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or a hydrazine radical according to formula 4.4,

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W_m and Y_m in formula 4 are absent or, independently of one another, are a carbonyl group,

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X_m is absent or is a hydrogen atom, a linear or branched C1-C30 alkylene radical, a linear, branched or cyclic C1-C30 alkenyl radical, a linear, branched or cyclic C1-C30 alkynylene radical, a linear, branched or cyclic C1-C30 alkoxy radical or a linear, branched or cyclic C1-C30 alkylamino radical containing primary amino groups.

z is optional and is a hydrogen atom, a halogen atom, a carbon atom, an oxygen atom, a sulfur atom, a nitrogen atom, an NH_3 -o group, a phosphorus atom, a methyl radical, a methenyl radical, a methynyl radical, a monovalent, divalent or trivalent to hexavalent diol radical, for example a diol radical of 2 to 14, preferably 4 to 10, carbon atoms, e.g. ethylene glycol, 1,3-propanediol, 1,10-decanediol, o-, m- or p-dihydroxycyclohexane, diethylene glycol, dipropylene glycol and preferably 1,4-butanediol, 1,6-hexanediol and bis(2-hydroxyethyl)hydroquinone, a triol radical, such as 1,2,4- or 1,3,5-trihydroxycyclohexane, glycerol and trimethylolpropane, or alcohols having a higher OH functionality, such as pentaerythritol or sucrose, or an amino alcohol or diamine radical, for example ethylenediamine, 1,2-propanediamine, 1,3-propanediamine, tetramethylenediamine, 2-methyl-1,5-diaminopentane, 1-methyl-2,4-diaminocyclohexane, 1,2-diaminocyclohexane, 1,3-diaminocyclohexane, 1,4-diaminocyclohexane or isophoronediamine.

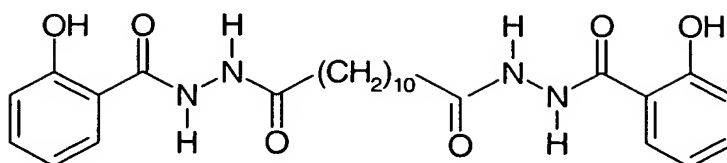
In formula 4, n is an integer from 1 to 6,
m is an integer from 1 to n and
o in the expression NH_3 -o is an integer from 1 to 3.

Examples of the preferred above structures are decamethylenedicarboxydisalicyloylhydrazides (ADK Stab CDA 6[®]) and 3-salicyloylamino-1,2,4-triazole (ADK Stab CDA 1[®]) according to formulae 5 and 6.

ADK Stab CDA 6

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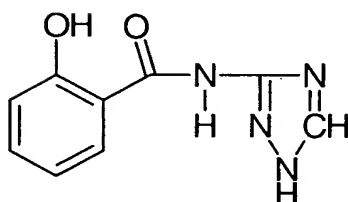
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ADK Stab CDA 1

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ADK Stab CDA 6 is particularly preferred.

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It may be advantageous to use mixtures of metal deactivators instead of a single metal deactivator. Mixtures which contain ADK Stab CDA 1 or ADK Stab CDA 6 are preferred, in particular those mixtures which contain ADK Stab CDA 6.

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In an embodiment of the present invention, the two stabilizing components (i) antioxidant and (ii) metal-deactivating component may be combined in one molecule. Examples of such a stabilizer are

35 2,2'-oxamidobis(ethyl-3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionates) and

N,N'-bis(3-(3',5'-di-tert-butyl-4'-hydroxyphenyl)propionylhydrazides).

40 For stabilizing thermoplastic polyurethanes, the components (i) and (ii) and, if required, (iii) are incorporated into the polyurethane.

In a preferred embodiment, the polyurethane to be stabilized

45 generally contains from 0.1 to 5% by weight, based on the total weight of the polyurethane, of antioxidant (i).

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If the TPU to be stabilized is a polyether-TPU having a Shore hardness of less than Shore 54 D, the antioxidant (i) is usually used in concentrations of from 0.1 to 5, preferably from 0.1 to 2, particularly preferably from 0.5 to 1.5, % by weight, based on 5 the total weight of the TPU.

If the TPU to be stabilized is a polyester-TPU or a polyether-TPU having a Shore hardness greater than or equal to Shore 54 D, the antioxidant (i) is usually used in concentrations of from 0.1 to 10 5, preferably from 0.1 to 1, particularly preferably from 0.2 to 0.75, % by weight, based on the total weight of the TPU.

The novel, stabilized TPU usually contains 0.01 - 2.5, preferably 0.05 - 1, particularly preferably 0.1 - 0.5, % by weight, based 15 on the total weight of the TPU, of the component (ii).

Where the components (i) and (ii) are added in the form of a compound which combines the two components, this compound is generally used in an amount of 0.01 - 2.5, preferably 0.05 - 1, 20 particularly preferably 0.1 - 0.5, % by weight, based on the total weight of the TPU. An antioxidant (component i) may then also be added, so that the concentration of component (i) corresponds to the preferred embodiment.

25 In a preferred embodiment, flameproofing agent, as component (iii) for reducing the flammability, may also be added to the TPU stabilized according to the invention and comprising the components (i) and (ii).

30 Suitable halogen-free flameproofing agents in addition to, for example, aluminum trihydrate and magnesium hydrate for particularly low-melting TPUs, are the triesters of phosphoric acid, such as trialkyl phosphates or triaryl phosphates, such as triphenyl phosphate. Oligomeric phosphoric esters or phosphonic 35 esters and cyclic phosphates which are derived from pentaerythritol, neopentylglycol or pyrocatechol are particularly preferred. These phosphoric esters can be used alone or as mixtures with one another or as mixtures with phosphonic esters. Usually, however, phosphoric esters or phosphonic esters are 40 used.

In a particularly suitable flameproofing combination, the phosphoric esters and/or phosphonic esters are used as mixtures together with one or more melamine derivatives for the TPU. In 45 this case, the weight ratio of phosphate or phosphonate to melamine derivative is preferably from 5:1 to 1:5. Suitable melamine derivatives are preferably melamine cyanurate, melamine

phosphate and melamine borate, particularly preferably melamine cyanurate.

However, it has surprisingly been found that in particular the
5 use of melamine derivatives without addition of phosphoric esters gives a particularly suitable TPU for the production of film-laminated flat conductors and cable sheaths, and this embodiment is therefore preferred.

10 A large number of chlorinated or brominated compounds may be used as halogen-containing flameproofing agents. Effective flameproofing agents are, for example, chlorinated polyethylene, chlorinated cyclopentadiene adducts, decabromodiphenyl oxide, decabromobiphenyl, poly(tetrabromobisphenol A glycidyl ether) and
15 poly(tetrabromobisphenol A carbonate) in combination with one another and/or with antimony(III) oxide as a synergistic agent and/or zinc borate. A number of other metal oxides, e.g. ZnO, B₂O₃, Fe₂O₃ or CaO, can also be added to the TPU in order to improve the flameproofing effect.

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Polytetrafluoroethylene and silica in very small amounts are suitable for reducing the tendency to drip flaming particles.

The amounts of the component (iii) which, if required, is added
25 for reducing the flammability vary greatly depending on the choice of the component, for example whether it is a halogen-containing or a halogen-free flameproofing agent. The additives (iii) of the preferred embodiment and the amounts in which they are used are described in EP-A-0617079 and
30 EP-A-0189130.

The flameproofing agents are added to the TPU in general in an amount of from 0.1 to 60, preferably from 1 to 40, particularly preferably from 5 to 25, % by weight, based on the total weight
35 of the stabilized TPU.

In a particularly preferred embodiment, melamine cyanurate is used as flameproofing component (iii), in an amount of 0.1 - 60, particularly preferably 5 - 40, in particular 15 - 25, % by
40 weight.

The components (i), (ii) and, if required, (iii) can be used for stabilizing elastomeric polyurethane, in particular thermoplastic polyurethanes, or for the preparation of stabilized
45 polyurethanes.

Thermoplastic polyurethane means that it is a polyurethane-based thermoplastic elastomer. A thermoplastic elastomer is an elastomer which remains thermoplastic when it is repeatedly heated and cooled in the temperature range typical for processing and use of the material. Thermoplastic is understood as meaning the property of a plastic of repeatedly softening at elevated temperature in a temperature range typical for it and of hardening on cooling and, in the softened state, being repeatedly moldable through flow as a molding, extrudate or shaped article to give semifinished products or articles.

Processes for the preparation of thermoplastic polyurethanes are generally known. In general, TPUs are prepared by reacting (a) isocyanates with (b) compounds reactive toward isocyanates and having a molecular weight (M_w) of from 500 to 10 000 and (c) chain extenders having a molecular weight of from 50 to 499, in the presence or absence of (d) catalysts and/or (e) conventional additives.

The starting components and processes for the preparation of the preferred polyurethanes are to be described below by way of example. The components (a), (b), (c) and, if required, (d) and/or (e) usually used in the preparation of the polyurethanes are to be described below by way of example:

- 25 a) Organic isocyanates (a) which may be used are generally known aliphatic, cycloaliphatic, araliphatic and/or aromatic isocyanates, for example tri-, tetra-, penta-, hexa-, hepta- and/or octamethylene diisocyanate, 2-methylpentamethylene 30 1,5-diisocyanate, 2-ethylbutylene 1,4-diisocyanate, pentamethylene 1,5-diisocyanate, butylene 1,4-diisocyanate, 1-isocyanato-3,3,5-trimethyl-5-isocyanatomethylcyclohexane (isophorone diisocyanate, IPDI), 1,4- and/or 1,3-bis(isocyanatomethyl)cyclohexane (HMDI), cyclohexane 35 1,4-diisocyanate, 1-methylcyclohexane 2,4- and/or 2,6-diisocyanate and/or dicyclohexylmethane 4,4'-, 2,4'- and 2,2'-diisocyanate, diphenylmethane 2,2'-, 2,4'- and/or 4,4'-diisocyanate (MDI), naphthylene 1,5-diisocyanate (NDI), tolylene 2,4- and/or 2,6-diisocyanate (TDI), diphenylmethane 40 diisocyanate, dimethylbiphenyl 3,3'-diisocyanate, diphenylethane 1,2-diisocyanate and/or phenylene diisocyanate.
- b) The generally known compounds reactive toward isocyanates may 45 be used as compounds (b) reactive toward isocyanates, for example polyesterols, polyetherols and/or polycarbonatediols, which are usually also referred to together by the term

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polyols, having molecular weights of from 500 to 8 000, preferably from 600 to 6 000, in particular from 800 to less than 3 000, and preferably having an average functionality of from 1.8 to 2.3, preferably from 1.9 to 2.2, in particular 2.

5 Polyetherpolyols, for example those based on generally known initiators and conventional alkylene oxides, for example ethylene oxide, propylene oxide and/or butylene oxide, are preferably used, particularly preferably polyetherols based on 1,2-propylene oxide and ethylene oxide and in particular

10 polyoxytetramethylene glycols. The polyetherols have the advantage that they have a higher stability to hydrolysis than polyesterols.

Other polyetherols which may be used are those having a low

15 degree of unsaturation. In the context of this invention, polyols having a low degree of unsaturation are understood as meaning in particular polyetheralcohols containing less than 0.02, preferably less than 0.01, meq/g of unsaturated compounds.

20 Such polyetheralcohols are generally prepared by an addition reaction of alkylene oxides, in particular ethylene oxide, propylene oxide or a mixture thereof, with the diols or triols described above, in the presence of highly active catalysts. Such highly active catalysts are, for example, cesium hydroxide and

25 multimetal cyanide catalysts, also referred to as DMC catalysts. A frequently used DMC catalyst is zinc hexacyanocobaltate. The DMC catalyst can be left in the polyetheralcohol after the reaction, but it is usually removed, for example by sedimentation or filtration.

30 Polybutadienediols having a molar mass of 500 - 10 000, preferably 1 000 - 5 000, in particular 2 000 - 3 000, g/mol may furthermore be used. TPUs prepared using these polyols can be radiation-crosslinked after thermoplastic processing. This leads,

35 for example, to better burning behavior.

Instead of one polyol, it is also possible to use mixtures of different polyols.

40 c) Chain extenders (c) used are generally known aliphatic, araliphatic, aromatic and/or cycloaliphatic compounds having a molecular weight of from 50 to 499, preferably difunctional compounds, for example diamines and/or alkanediols having 2 to 10 carbon atoms in the alkylene radical, in particular

45 1,4-butanediol, 1,6-hexanediol and/or di-, tri-, tetra-, penta-, hexa-, hepta-, octa-, nona- and/or decaalkylene glycols of 3 to 8 carbon atoms, preferably corresponding

oligo- and/or polypropylene glycols, it also being possible to use mixtures of the chain extenders.

The components a) to c) are particularly preferably difunctional compounds, i.e. diisocyanates (a), difunctional polyols, preferably polyetherols (b) and difunctional chain extenders, preferably diols.

- d) Suitable catalysts which accelerate in particular the reaction between the NCO groups of the diisocyanates (a) and the hydroxyl groups of the components (b) and (c) are the conventional tertiary amines known from the prior art, e.g. triethylamine, dimethylcyclohexylamine, N-methylmorpholine, N,N'-dimethylpiperazine, 2-(dimethylaminoethoxy)ethanol, diazabicyclo[2.2.2]octane and the like, and in particular organic metal compounds, such as titanate esters, iron compounds, e.g. iron(III) acetylacetonate, tin compounds, e.g. tin diacetate, tin dioctanoate, tin dilaurate or the dialkyltin salts of aliphatic carboxylic acids, such as dibutyltin diacetate, dibutyltin dilaurate or the like. The catalysts are usually used in amounts of from 0.0001 to 0.1 part by weight per 100 parts by weight of polyhydroxy compound (b).
- e) In addition to catalysts (d), conventional assistants and/or additives (e) may also be added to the components (a) to (c). Examples are blowing agents, surface-active substances, fillers, nucleating agents, lubricants and mold release agents, dyes and pigments, if required further antioxidants which can be used in addition to the component (i), for example against hydrolysis, light, heat or discoloration, inorganic and/or organic fillers, reinforcing agents and plasticizers. In a preferred embodiment, the component (e) also includes hydrolysis stabilizers, for example polymeric and low molecular weight carbodiimides.

In addition to said components a), b) and c) and, if required, d) and e), chain regulators, usually having a molecular weight of from 31 to 499, may also be used. Such chain regulators are compounds which have only one functional group reactive toward isocyanates, e.g. monofunctional alcohols, monofunctional amines and/or monofunctional polyols. By means of such chain regulators, flow behavior, in particular in the case of TPUs, can be established in a controlled manner. Chain regulators can be used in general in an amount of from 0 to 5 parts by weight, preferably from 0.1 to 1 part by weight, based on 100 parts by

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weight of the component b), and by definition are included in the component c).

All molecular weights stated in this document have the unit
5 [g/mol].

For establishing hardness of the TPUs, the components (b) and (c) can be varied within relatively wide molar ratios. Molar ratios of from 10 : 1 to 1 : 10, in particular from 1 : 1 to 1 : 4, for
10 component (b) to the total amount of chain extenders (c) to be used have proven useful, the hardness of the TPU increasing with increasing content of (c).

The reaction can be carried out at customary indices, preferably
15 at an index of from 60 to 120, particularly preferably from 80 to 110. The index is defined by the ratio of the total amount of isocyanate groups of component (a) which are used in the reaction to those groups of the components (b) and (c) which are reactive toward isocyanates, i.e. the active hydrogen atoms. At an index
20 of 100, there is one active hydrogen atom of the components (b) and (c), i.e. one function reactive toward isocyanates, per isocyanate group of the component (a). At indices above 100, more isocyanate groups are present than OH groups.

25 The TPU can be prepared by the known processes, either continuously, for example using reaction extruders or the belt process in the one-shot or prepolymer process, or batchwise by the known prepolymer process. In these processes, the components (a), (b), (c) and, if required, (d) and/or (e) which are reacted
30 can be mixed with one another in succession or simultaneously, the reaction starting immediately.

In the extruder process, the components (a), (b), (c) and, if required, (d) and/or (e) are fed individually or as a mixture
35 into the extruder, for example at from 100 to 280°C, preferably from 140 to 250°C, and are reacted. The TPU obtained is extruded, cooled and granulated.

After the synthesis, the TPU can, if required, be modified by
40 compounding in an extruder. By means of this compounding, the TPU can be modified, for example, in its melt index or its granular form to meet the requirements.

The components (i) - (iii) can be metered in during the synthesis or the processing of the TPU. Concentrates which contain the components (i) and/or (ii) and/or (iii) can also be prepared and metered into the TPU during the processing.

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The case in which the TPU is stabilized during the synthesis or compounding with component i) and, if required, component iii) and a concentrate of the component ii) is metered into the TPU during the processing is preferred.

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Such a novel metal deactivator concentrate contains in general a TPU, preferably a polyether-TPU, and from more than 2.5 to 40, preferably from 3 to 20, particularly preferably from 4 to 10, % by weight, based on the total weight of the thermoplastic

15 polyurethane, of metal deactivator (ii). In addition, the concentrate can, if required, also contain an antioxidant, preferably a phenolic antioxidant, usually in an amount of from 0.1 to 5, preferably from 0.5 to 2, % by weight.

20 The novel concentrate can be prepared by adding the metal deactivator (ii) and, if required, the antioxidant (i) to the TPU during the synthesis, or by melting a TPU in a batch kneader or in an extruder and adding the metal deactivator and, if required, the antioxidant to the TPU during the synthesis. The addition in
25 an extruder, in particular a twin-screw extruder, is preferred.

The amount of this concentrate is preferably chosen so that the concentration of the component ii) in the TPU thus treated corresponds to the particularly preferred concentration for the
30 component ii) in the TPU. Preferred concentrates are those which are metered into the TPU in an amount of 0.5 - 10, preferably 1 - 4, % by weight, based on the total amount of the TPU.

The processing of the TPUs prepared according to the invention
35 which are usually present in the form of granules or powder, to give injection molded and extruded articles, for example the desired films, shaped articles, rollers, fibers, claddings in automobiles, hoses, cable connectors, bellows, trailing cables, cable sheaths, seals, belts or damping elements, is effected by
40 conventional methods, for example injection molding or extrusion. Such injection molded or extruded articles may also consist of compounds containing the novel TPU and at least one further thermoplastic, in particular a polyolefin, polyester, polyether, polystyrene or styrene copolymer or polyoxymethylene. In
45 particular, TPU prepared according to the invention can be used

for the production of film-laminated flat conductors and for the sheathing of copper cables.

Film-laminated flat conductors are generally understood as
5 meaning flat, flexible conductor strips which are composed of one or more, preferably parallel, metal strips which are incorporated between two films of plastic so that an insulation is produced. The dimensions of the film-laminated flat conductors are not limited. Usually, they have a length of from 5 cm to 100 m and a
10 width of from 0.5 cm to 30 cm, depending on their use.

According to the present invention, the plastics films contain thermoplastic polyurethane. They preferably comprise thermoplastic polyurethane, particularly preferably the TPU
15 stabilized according to the invention. The films generally have a thickness of from 1 μm to 5 mm, preferably from 10 μm to 1 mm, particularly preferably from 25 μm to 250 μm . The adhesion between films is usually achieved by heating and pressing and/or by the use of adhesives.

20 A suitable metal strip is in general any conductive material. However, metals such as iron, zinc, copper or alloys thereof are preferably used, particularly preferably copper. The metal strips generally have a thickness of from 1 to 500 μm , preferably from 10
25 to 250 μm , particularly preferably from 30 to 100 μm .

A plurality of processes are possible for the production of the novel film-laminated flat conductors. In a preferred embodiment, first two TPU films preferably coated on one side with adhesive
30 are provided. Metal strips, for example 30 - 100 μm thick copper strips, are then incorporated or laminated between these two films. A preferred adhesive is a resistant adhesive.

In a further preferred embodiment, the film laminate is produced
35 in one operation, by coextrusion of the TPU films with metal strips, preferably with the copper strips. The additional processing step of lamination is thus dispensed with.

It is furthermore possible to produce the novel film-laminated
40 flat conductors by pressing the copper conductor into the TPU film.

The novel film-laminated flat conductors have a wide range of applications. They are preferably used as flexible conductors in
45 means of transport, electrical articles and machines. Examples of suitable means of transport are motor vehicles, for example cars or trucks, railroad vehicles, aircraft or ships. Examples of

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electrical articles are household appliances, televisions, stereo systems, video recorders, computers and accessories, printers and accessories, copiers and accessories, scanners and accessories, switch cabinets and control systems. Examples of machines are
5 packaging machines, robots, cutting machines, machine tools, injection molding machines, extruders, calenders, film blowing machines, CAD machines, milling units, punches, presses, turning machines, construction machines, e.g. diggers, wheel loaders, cranes, conveyor systems, industrial trucks, sorting machines,
10 assembly lines, process monitoring systems and process control stations.

The examples which follow illustrate the invention.

15 Example 1

Example 1 describes the preparation of a novel stabilized TPU by the manual casting method.

20 In order to simulate the large surface areas as exist in film-laminated flat conductors, copper powder was incorporated directly.

1 000 g of a polytetrahydrofuran having a molar mass of 1 000
25 g/mol (PTHF 1000) were heated to about 90°C in a tinplate bucket. Antioxidant (component i; for amount see table 1) and 125 g of butanediol were then added while stirring. In addition, the amounts of copper powder (particle size 0.04 mm) and metal deactivator (ii) stated in table 1 were metered in. The solution
30 was heated to 80°C while stirring. 600 g of 4,4'-MDI were then added and stirring was carried out until the solution was homogeneous. Thereafter, the TPU was poured into a shallow dish and heated first for 10 minutes at 125°C on a hotplate and then for 15 hours at 110°C in an oven.

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Experiment number	Antioxidant 1	Amount	Antioxidant 2	Amount	Metal deactivator	Amount	Cu
5 1.1	Irganox [®] 1010	8g	Irganox [®] 1098	8g	ADK Stab [®] CDA 1	1.75 g	5.2 g
1.2	Irganox [®] 1010	8g	Irganox [®] 1098	8g	ADK Stab [®] CDA 6	1.75 g	5.2 g
1.3	Irganox [®] 1010	8g	Irganox [®] 1098	8g	Irganox [®] MD 1024	1.75 g	5.2 g
10 1.4	Irganox [®] 1010	8g	Irganox [®] 1098	8g	Eastman [®] Inhibitor	1.75 g	5.2 g
1.5	Irganox [®] 1010	8g	Irganox [®] 1098	8g	Palmarole [®] MDA P10	1.75 g	5.2 g
15 1.6	Irganox [®] 1010	8g	Irganox [®] 1098	8g	Palmarole [®] MDA P11	1.75 g	5.2 g
1.7	Irganox [®] 1010	8g	Irganox [®] 1098	8g	Hostanox [®] OSP 1	1.75 g	5.2 g
1.8	Irganox 1010	8g	Irganox [®] 1098	8g	—	—	5.2 g

20 Table 1

Irganox[®] and Tinuvin[®] are trade names of Ciba Spezialitätenchemie GmbH, Lampertsheim, Germany

25 ADK[®] Stab and Palmarole[®] MDA are trade names of Asahi Denka Kogyo K.K., Tokyo, Japan.

Eastman Inhibitor[®] is a trade name of Eastman Chemical Company, Kingsport, TN, USA

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Hostanox[®] is a trade name of Clariant GmbH, Germany.

Example 2

35 Example 2 describes the production of test plates and the aging tests

The cast slabs from example 1 were granulated and injection molded to give 2 mm thick sheets. S 2 test specimens were punched
40 out of these injection-molded sheets. These test specimens were aged in an oven at 130°C. The test specimens were removed from the oven at predefined intervals and tested according to DIN 53504. Figure 1 shows the curve for the elongation at break for the samples according to examples 1.1 to 1.8. It is clear that the
45 addition of metal deactivators improves the stability of the TPU.

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Moreover, it is found that the preferred metal deactivators from experiments 1.1 and 1.2 have a particularly good effect.

Example 3

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Example 3 describes the preparation of a metal deactivator concentrate

1 000 g of PTHF 1000 were heated to about 90°C in a tinplate
10 bucket. 8 g of Irganox® 1010 and 8 g of Irganox® 1098 and 125 g of butanediol were then added while stirring. In addition, 50 g of ADK Stab CDA 1® (experiment 3.1) and 50 g of ADK Stab CDA 6® (experiment 3.2) were metered in. The solution was heated to 80°C while stirring. 600 g of 4,4'-MDI were then added and stirring
15 was carried out until the solution was homogeneous. Thereafter, the TPU was poured into a shallow dish and heated first for 10 minutes at 125°C on a hotplate and then for 15 hours at 110°C in an oven.

20 Example 4

Example 4 describes the production of a TPU film with the aid of a metal deactivator concentrate.

25 4% of a concentrate from example 3 were added to various polyether-TPU of the Elastollan® series and the granules were thoroughly mixed and then processed on a Dr. Collin Cill-Roll unit 136-350 and a 250 mm slot die. Experiments in which the novel stabilized TPU was not used are indicated by C. A film
30 thickness of 50 µm was established. Both concentrates mentioned in example 3 can be readily incorporated. Table 2 gives information about the experiments carried out.

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Experiment	TPU	Component (i)	Amount	Concentrate	Amount	Component (iii)
5	C 4.1	1154 D	Irganox [®] 1125	0.5% by wt.	—	—
	4.2	1154 D	Irganox [®] 1125	0.5% by wt.	3.1	4% by wt.
	4.3	1154 D	Irganox [®] 1125	0.5% by wt.	3.2	4% by wt.
	C 4.4	1185 A	Irganox [®] 1125	1% by wt.	—	—
	4.5	1185 A	Irganox [®] 1125	1% by wt.	3.1	4% by wt.
10	4.6	1185 A	Irganox [®] 1125	1% by wt.	3.2	4% by wt.
	C 4.7	1154D	Irganox [®] 1125	0.5% by wt.	—	—
	FHF					yes
	4.8	1154 D	Irganox [®] 1125	0.5% by wt.	3.1	4% by wt.
15	FHF					yes
	4.9	1154 D	Irganox [®] 1125	0.5% by wt.	3.2	4% by wt.
	FHF					yes
20	C 4.10	1185 A	Irganox [®] 1125	1% by wt.	—	—
	FHF					yes
	4.11	1185 A	Irganox [®] 1125	1% by wt.	3.1	4% by wt.
	FHF					yes
	4.12	1185 A	Irganox [®] 1125	1% by wt.	3.2	4% by wt.
	FHF					yes

Table 2

25 Elastollan[®] is the trade name of Elastogran GmbH, Lemförde, Germany.

Example 5

30 Production of film-laminated flat conductors

Films whose production was described in example 4 were pressed between two copper sheets and heated at 110°C in an oven. After 250 hours, the films were removed. Table 3 shows the extent to which the individual films have become discolored.

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	Experiment	TPU film	Discoloration
	C 5.1	4.1	Strongly yellowed
	5.2	4.2	Slightly yellowed
	5.3	4.3	Slightly yellowed
5	C 5.4	4.4	Strongly yellowed
	5.5	4.5	Slightly yellowed
	5.6	4.6	Slightly yellowed
	C 5.7	4.7	Strongly yellowed
	5.8	4.8	Slightly yellowed
10	5.9	4.9	Slightly yellowed
	C 5.10	4.10	Strongly yellowed
	5.11	4.11	Slightly yellowed
	5.12	4.12	Slightly yellowed

15 Table 3

It is clear that the films from the examples which contain no metal deactivators have a stronger yellow discoloration.

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